

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>7</sup> :</b> <b>C10L 1/08</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 00/11117</b> <b>(43) International Publication Date:</b> 2 March 2000 (02.03.00)
<b>(21) International Application Number:</b> PCT/US99/17013 <b>(22) International Filing Date:</b> 27 July 1999 (27.07.99)  <b>(30) Priority Data:</b> 09/138,130                      21 August 1998 (21.08.98)                      US  <b>(71) Applicant:</b> EXXON RESEARCH AND ENGINEERING COMPANY [US/US]; P.O. Box 390, 180 Park Avenue, Florham Park, NJ 07932-0390 (US).  <b>(72) Inventors:</b> BERLOWITZ, Paul, Joseph; 4 Nicole Terrace, Glen Gardner, NJ 08826 (US). WITTENBRINK, Robert, Jay; 6018 Riverchase Trail, Kingwood, TX 77345 (US). COOK, Bruce, Randall; 1101 Monroe Drive, Stewartville, NJ 08886 (US).  <b>(74) Agents:</b> SIMON, Jay et al.; Exxon Research and Engineering Company, P.O. Box 390, Florham Park, NJ 07932-0390 (US).		<b>(81) Designated States:</b> BR, CA, JP, NO, SG, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> IMPROVED STABILITY FISCHER-TROPSCH DIESEL FUEL AND A PROCESS FOR ITS PRODUCTION  <b>(57) Abstract</b>  A Fischer-Tropsch derived distillate fraction is blended with either a raw virgin condensate fraction or a mildly hydrotreated virgin condensate to obtain a stable inhibited distillate fuel.		

*FOR THE PURPOSES OF INFORMATION ONLY*

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

IMPROVED STABILITY FISCHER-TROPSCH DIESEL FUEL AND A  
PROCESS FOR ITS PRODUCTION

## FIELD OF THE INVENTION

This invention relates to stable, inhibited middle distillates and their preparation. More particularly, this invention relates to stable, inhibited middle distillates, useful as fuels e.g., kerosene, diesel, or as fuel blending components, in which a Fischer-Tropsch derived distillate is blended with a virgin distillate.

## BACKGROUND OF THE INVENTION

Distillate fuels derived from Fischer-Tropsch processes are often hydrotreated to eliminate unsaturated materials, e.g., olefins, and most, if not all, oxygenates. The hydrotreating step is often combined with mild hydroisomerization resulting in the formation of iso-paraffins, often necessary for meeting pour point specifications for distillate fuels, particularly fuels heavier than gasoline, e.g., diesel and jet fuels.

Fischer-Tropsch distillates, by their nature, have essentially nil sulfur and nitrogen, these elements having been removed upstream of the Fischer-Tropsch reaction because they are poisons, even in rather small amounts, for known Fischer-Tropsch catalysts. As a consequence, Fischer-Tropsch derived distillate fuels are inherently stable, the compounds that may lead to instability, e.g., by oxidation, having been removed either upstream of the reaction or downstream in subsequent hydrotreating steps. While stable, these distillates have no inherent inhibitors for maintaining oxidative stability. Thus, upon the onset of oxidation, as in the formation of peroxides, a measure of oxidative stability, the distillate

- 2 -

has no inherent mechanism for inhibiting oxidation. These materials may be viewed as having a relatively long induction period for oxidation, but upon initiation of oxidation, the material efficiently propagates oxidation.

Virgin distillates as may be obtained from conventional petroleum sources are usually a constituent of distillate fuels, and contain sulfur in varying concentrations. The addition, usually small amounts, of virgin distillate to Fischer-Tropsch distillates provides a facile method for stabilizing Fischer-Tropsch derived fuels against oxidation.

## SUMMARY OF THE INVENTION

In accordance with this invention, a blended middle distillate, useful as a fuel or a fuel blending component, and having both stability and resistance to oxidation comprises: a Fischer-Tropsch (F-T) derived distillate and a virgin distillate fraction, and wherein the sulfur content of the blend is  $\geq 1$  ppm by wt.

## BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows the effect on peroxide number of adding 1%, 5%, and 25% by weight of a virgin distillate to a Fischer-Tropsch derived distillate fuel.

Figure 2 shows the effect on peroxide number of adding a mildly hydrotreated virgin distillate having 210 ppm sulfur in amounts of 0.1, 0.5, 5.0, and 25% by weight to a Fischer-Tropsch derived fuel.

- 3 -

In each figure the peroxide number after 28 days is shown on the ordinate and the weight fraction Fischer-Tropsch derived fuel is shown on the abscissa.

In the absence of any known effects on the addition of a relatively less stable fuel with a relatively more stable, but uninhibited fuel, one would expect the peroxide number to fall on a straight line connecting the peroxide numbers for a 100% F-T derived fuel and a 100% virgin distillate fuel, shown in the drawings as a dotted line.

The data in the drawings make it abundantly clear that small amounts of virgin distillate, when added to a Fischer-Tropsch derived fuel can, and do, have a significant effect on the long term stability of the F-T derived fuel.

The distillate fraction for either the Fischer-Tropsch derived material or the gas field condensate is a C<sub>8</sub>-700°F stream, preferably comprised of a 250-700°F fraction, and preferably in the case of diesel fuels or diesel range fuels, a 320-700°F fraction.

The virgin distillate is preferably a distillate fraction that is essentially untreated, or stated otherwise, is in the substantial absence of any treatment materially changing the boiling point of the hydrocarbon liquids in the virgin distillate. Thus, the condensate has not been subjected to conversion by means that may significantly or materially change the boiling point of the liquid hydrocarbons in the virgin distillate. The virgin distillate, however, may have been de-watered, desalted, distilled to the proper fraction, or mildly hydrotreated, none of which significantly effects the boiling point of the liquid hydrocarbons of the virgin distillate.

In one embodiment, the virgin distillate may be subjected to hydrotreating, e.g., mild hydrotreating, that reduces sulfur content and olefinic content, but does not significantly or materially effect the boiling point of the liquid hydrocarbons. Thus, hydrotreating, even mild hydrotreating is usually effected in the presence of a catalyst, such as supported Co/Mo, and some hydrocracking may occur. In the context of this invention, unprocessed virgin distillate includes virgin distillate subjected to mild hydrotreating which is defined as hydrotreating that does not materially change the boiling point of the liquid hydrocarbons and maintains sulfur levels of  $>10$  ppm, preferably  $\geq 20$  ppm, more preferably  $\geq 30$  ppm, still more preferably  $\geq 50$  ppm. Thus, the forms of sulfur that act as oxidation inhibitors are not present in sufficient concentrations in the virgin distillate to provide inhibiting effects.

The result of this mixture is a distillate fraction, preferably a 250-700°F fraction and more preferably a 320-700°F that is both stable and resistant to oxidation. Oxidation stability is often determined as a build up of peroxides in the sample under consideration. While there is no standard for the peroxide content of fuels, there is general acceptance that stable fuels have a peroxide number of less than about 5, preferably less than about 4, and desirably less than about 1.

The Fischer-Tropsch process is well known and preferably utilizes a non-shifting catalyst such as cobalt or ruthenium or mixtures thereof, preferably cobalt, and more preferably a promoted cobalt, particularly where the promoter is rhenium. Such catalysts are well known and described in U.S. Patents 4,568,663 and 5,545,674.

Non-shifting Fischer-Tropsch reactions are well known and may be characterized by conditions that minimize the formation of CO<sub>2</sub> by-products. These conditions can be achieved by a variety of methods, including one or more of the following: operating at relatively low CO partial pressures, that is, operating at hydrogen to CO ratios of at least about 1.7/1, preferably about 1.7/1 to 2.5/1, more preferably at least about 1.9/1 and in the range 1.9/1 to about 2.3/1, all with an alpha of at least about 0.88, preferably at least about 0.91; temperatures of about 175° - 240°C, preferably about 180°C - 220°C, using catalysts comprising cobalt or ruthenium as the primary Fischer-Tropsch catalysis agent. A preferred process for conducting the Fischer-Tropsch process is described in U.S. Patent 5,348,982.

The products of the Fischer-Tropsch process are primarily paraffinic hydrocarbons, although very small amounts of olefins, oxygenates, and aromatics may also be produced. Ruthenium catalysts produce paraffins primarily boiling in the distillate range, i.e., C<sub>10</sub>-C<sub>20</sub>; while cobalt catalysts generally produce more heavier hydrocarbons, e.g., C<sub>20</sub><sup>+</sup>.

The diesel fuels produced from Fischer-Tropsch materials generally have high cetane numbers, usually 50 or higher, preferably at least 60, and more preferably at least about 65.

Virgin distillates may vary in composition from field to field, but the virgin distillates will have some similar characteristics, such as: a boiling range of 250-700°F, preferably 320-700°F, derived from petroleum sources. Virgin middle distillates are always a mixture of paraffins, naphthene and aromatic hydrocarbons, as well as organic sulfur and nitrogen compounds. The exact amounts of each of these species is widely variable, but in most cases paraffins

- 6 -

range from 20-70%, naphthas 10-40% and aromatic from 5-40%. Sulfur can range from a few hundred ppm to several percent.

The F-T derived middle distillate and the virgin middle distillate may be mixed in wide proportions, and as shown above, small fractions of virgin distillate can significantly effect the peroxide number of the blend. Thus, blends of 1-50 wt% virgin distillate with 99-50 wt% F-T derived distillate may readily be formed. Preferably, however, the virgin distillate is blended at levels of 1-40 wt% with the F-T derived distillate, more preferably 1-30 wt%.

The stable middle distillate blend of F-T derived distillate and virgin distillate may then be used as a fuel, e.g., diesel or jet, and preferably a fuel heavier than gasoline, or the blend may be used to upgrade or volume enhance petroleum based fuels. For example, a few percent of the blend can be added to a conventional petroleum based fuel for enhancing cetane number, typically 2-20%, preferably 5-15%, more preferably 5-10%; alternatively, greater amounts of the blend can be added to the petroleum based fuel to reduce sulfur content of the resulting blend, e.g., about 30-70%. Preferably, the blend of this invention is mixed with fuels having low cetane numbers, such as less than 50, preferably less than 45.

The blend of virgin distillate and Fischer-Tropsch distillate will preferably have a sulfur content of at least 2 ppm by weight; more preferably at least about 5 ppm, still more preferably at least about 15 ppm, still more preferably about  $\geq 25$  ppm, and yet more preferably  $\geq 50$  ppm. The blend may contain up to about 250 ppm S, preferably less than about 200 ppm S, more preferably less than 100 ppm S, still more preferably less than 50 ppm, and yet more preferably less than 30 ppm S.



Fischer-Tropsch derived distillates useful as fuels can be obtained in a variety of ways known to those skilled in the art, e.g., in accordance with the procedures shown in U.S. Patent 5,689,031 or allowed U.S. application S.N. 798,376, filed.

Additionally, many papers have been published in which F/T derived distillate fuels are obtained by hydrotreating/hydroisomerizing all or appropriate fractions of Fischer-Tropsch process products and distilling the treated/isomerized product to the preferred distillate fraction.

Fischer-Tropsch distillates useful as fuels or fuel blending components are generally characterized as being:

>80 wt%, preferably >90 wt%, more preferably >95 wt% paraffins, having an iso/normal ratio of 0.1 to 10, preferably 0.3 to 3.0, more preferably 0.7 to 2.0; sulfur and nitrogen of less than 1 ppm each, preferably less than 0.5, more preferably less than 0.1 ppm each;  $\leq 0.5$  wt% unsaturates (olefins and aromatics), preferably  $\leq 0.1$  wt%; and less than 0.5 wt% oxygen on a water free basis, preferably less than about 0.3 wt% oxygen, more preferably less than 0.1 wt% oxygen and most preferably nil oxygen. (The F-T distillate is essentially free of acids.)

- 8 -

The iso paraffins of a F-T derived distillate are mono-methyl branched, preferably primarily mono methyl branched and contain exceedingly small amounts of cyclic paraffins, e.g., cyclo hexanes. Preferably, the cyclic paraffins of the F-T distillate are not readily detectable by standard methods, e.g., gas chromatography.

The following examples serve to illustrate but not limit the invention:

Example 1: Stability of Fischer-Tropsch derived Distillate fuels: Blends with raw virgin distillate

A Fischer-Tropsch fuel produced by the process described in U.S. Serial No. 544,343) was distilled to a nominal 250-700°F boiling point encompassing the distillate range. This material was tested according to a standard procedure for measuring the buildup of peroxides: First a 4 oz. sample was placed in a brown bottle and aerated for 3 minutes. An aliquot of the sample is then tested according to ASTM D3703-92 for peroxides. The sample is then capped and placed into a 60°C oven for 1 week. After this time the peroxide number is repeated, and the sample is returned to the oven. The procedure continues each week until 4 weeks have elapsed and the final peroxide number is obtained. A value of <1 is considered a stable, distillate fuel.

This fuel was blended with a raw virgin distillate material in amount ranging from 0.1 to 25% to determine the effect on the final peroxide number. The data is shown in the Table 1 below

- 9 -

Table 1

% F-T Fuel	% Virgin Condensate	Initial Peroxide #	Final Peroxide #	S <sub>1</sub> ppm in Blend
100	0	0	24.06	0
75	25	0	0.63	550
95	5	0	0.68	110
99	1	0	0.88	21
99.9	0.1	0	13.17	2
0	100	0	0	2100

There is a significant effect of 0.1% of the raw virgin distillate which reduced the peroxide number close to 50%, occurring at a sulfur level of only 2 ppm in the blend (2100 ppm in the raw virgin distillate neat).

Example 2: Stability of Fischer-Tropsch derived Distillate fuels: Blends with hydrotreated virgin distillate

A Fischer-Tropsch fuel produced by the same (as in example 1) was distilled to a nominal 250-700°F boiling point encompassing the distillate range. This material was tested according to a standard procedure as described in Example 1.

This fuel was blended with a virgin distillate material which had been conventionally hydrotreated to 290 ppm S. Blends were in amounts ranging from 0.1 to 25% to determine the effect on the final peroxide number. The data is shown in Table 2, below:

- 10 -

Table 2

% F-T Fuel	% Virgin Condensate	Initial Peroxide #	Final Peroxide #	S <sub>1</sub> ppm in Blend
100	0	0	24.06	0
75	25	0	0.84	73
95	5	0	3.87	15
99	1	0	9.47	3
99.9	0.1	0	25.26	0.3
0	100	0	0	2100

As in Example 1, a significant benefit can be obtained at low sulfur concentrations. At a concentration of only 1% virgin distillate (3 ppm S in the blend), the buildup of peroxides is reduced 61%. In another test, at 0.3 ppm S or 0.1% hydrotreated condensate there is no significant effect, and the results for the neat F-T fuel are reproduced to within 5%.

These results indicate that a virgin distillate stream blended with an F-T fuel which has at least 2 ppm S in the final blend will substantially inhibit peroxide growth in the final fuel. The virgin distillate may be hydrotreated to remove 90% or more of the original S in the petroleum and still function effectively.

- 11 -

## CLAIMS:

1. A blend material useful as a distillate fuel or as a blending component for a distillate fuel comprising: (a) a Fischer-Tropsch derived distillate comprising a C<sub>8</sub>-700°F fraction, and (b) a virgin distillate comprising a C<sub>8</sub>-700°F fraction, wherein the sulfur content of the blend material is  $\geq 2$  ppm by wt.
2. The blend material of claim 1 wherein the Fischer-Tropsch distillate is a 250-700°F fraction and has a sulfur content of less than 1 ppm by wt.
3. The blend material of claim 1 wherein the virgin distillate is selected from the group consisting of raw virgin distillate and mildly hydrotreated virgin distillate where the boiling range of the distillate is not materially changed.
4. The blend material of claim 3 wherein the sulfur content of the virgin distillate is  $\geq 10$  ppm.
5. The blend material of claim 3 wherein the proportion of (a) to (b) is about 99/1 to 50/50.
6. The blend material of claim 5 wherein the proportion of (b) in the blend ranges from about 1-40%.
7. The blend material of claim 5 wherein the proportion of (b) in the blend ranges from about 1-30%.
8. The blend material of claim 6 further blended with a petroleum derived distillate.

- 12 -

9. The further blend of claim 8 wherein the blend material is about 30-70% of the further blend.

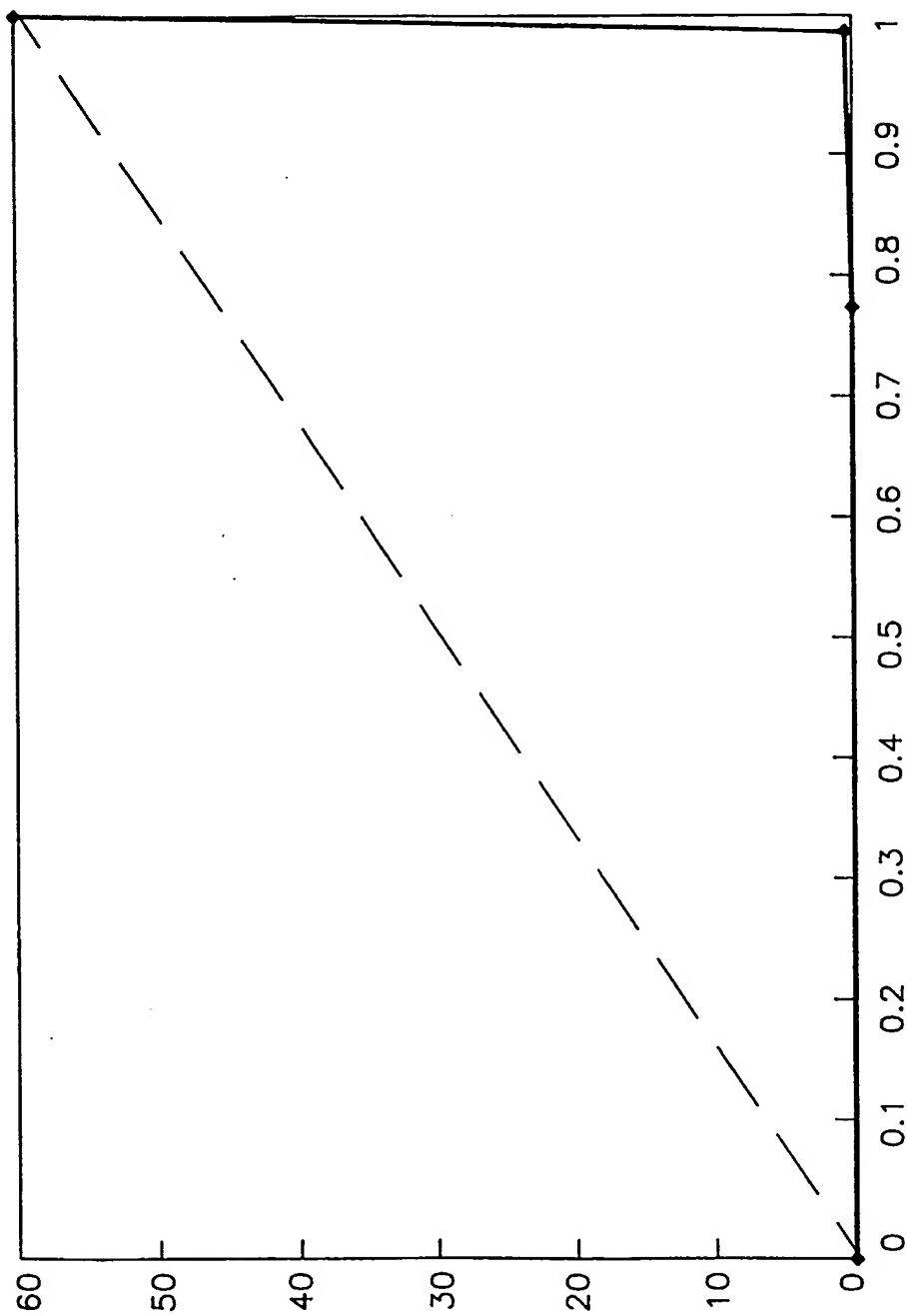


FIG. 1

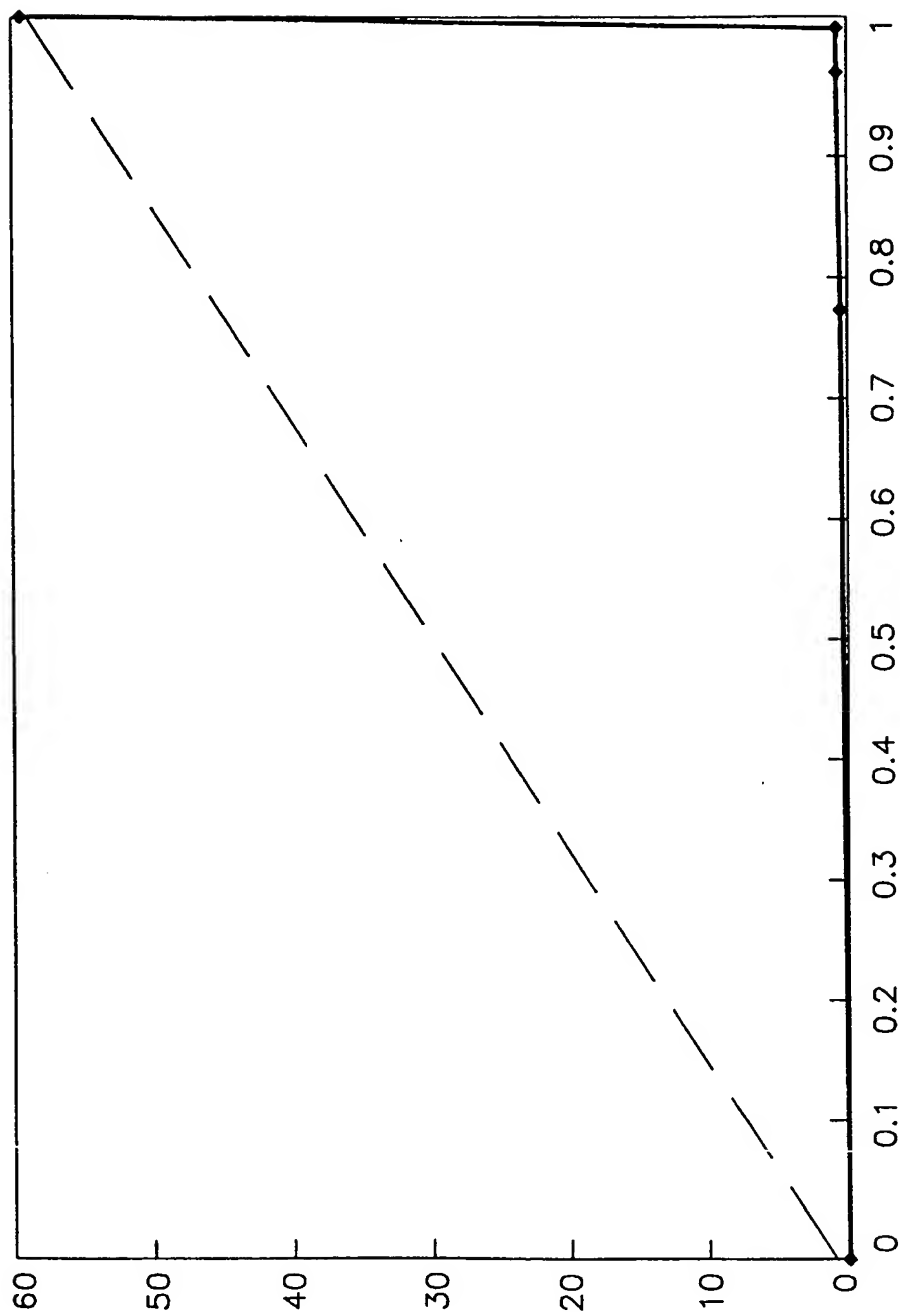


FIG. 2



# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/17013

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C10L1/08

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C10L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 98 34998 A (EXXON RESEARCH ENGINEERING CO) 13 August 1998 (1998-08-13) claims 1,4-6,10 ---	1,3,5
A	WO 97 14769 A (EXXON RESEARCH ENGINEERING CO) 24 April 1997 (1997-04-24) claims 1,12 ---	1
A	WO 96 26994 A (EXXON CHEMICAL PATENTS INC ;FAVA CARLOS S (GB); CAPROTTI RINALDO ( ) 6 September 1996 (1996-09-06) claims 1,2,4,5 -----	1-7

☐

Further documents are listed in the continuation of box C.

☒

Patent family members are listed in annex.

### \* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

27 October 1999

Date of mailing of the international search report

10/11/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

De Herdt, O

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/17013

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9834998 A	13-08-1998	US 5814109 A ZA 9800621 A	29-09-1998 22-07-1998
WO 9714769 A	24-04-1997	AU 7395196 A CA 2229433 A CN 1197476 A EP 0885275 A NO 981712 A	07-05-1997 24-04-1997 28-10-1998 23-12-1998 16-04-1998
WO 9626994 A	06-09-1996	CA 2213656 A DE 69600598 D DE 69600598 T EP 0812346 A JP 11501063 T	06-09-1996 08-10-1998 28-01-1999 17-12-1997 26-01-1999